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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT) WO 97/47708 (51) International Patent Classification 6: (11) International Publication Number: **A1** C10G 27/12, A01N 37/00 18 December 1997 (18.12.97) (43) International Publication Date: (81) Designated States: AU, AZ, BR, CA, CN, DE, DK, ES, GB, PCT/GB97/01537 (21) International Application Number: GE, KR, KZ, MX, NO, NZ, RU, TM, TR, UA, US, UZ, VN, Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, 9 June 1997 (09.06.97) (22) International Filing Date: TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). (30) Priority Data: 8 June 1996 (08.06.96) GB 9612035.7 **Published** With international search report. (71) Applicant (for all designated States except US): OILFIELD Before the expiration of the time limit for amending the CHEMICAL ADDITIVES LTD. [GB/GB]; Craigshaw claims and to be republished in the event of the receipt of Road, West Tullos, Aberdeen AB1 3AP (GB). amendments. (72) Inventor; and (75) Inventor/Applicant (for US only): BRANKLING, David [GB/GB]; 18 Corse Gardens, Kingswells, Aberdeen AB15 (74) Agent: MURGITROYD & COMPANY; 373 Scotland Street, Glasgow G5 8QA (GB).

(54) Title: METHOD OF INHIBITING RESERVOIR SOURING BY BACTERIA

(57) Abstract

A method of inhibiting the growth of bacteria, especially anaerobic sulphate reducing bacteria, within oil and gas reservoirs by the addition of peroxy compounds to injected flood water is described. Suitable peroxy compounds include one or more of methyl ethyl ketone peroxide, cyclohexanone peroxide, acetyl acetone peroxide, diacetone alcohol peroxide, dibenzoyl peroxide, ditertiary butyl peroxide, tertiary butyl peroxide, hydrogen peroxide, tertiary butyl peroxy benzoate, inorganic and organic peroxycarbonates, or functional equivalents or derivatives of these compounds. The peroxy compound additives have the advantage of being thermally sensitive such that penetration into the reservoir and placement in strategic zones can be achieved leading to sterilisation deep within the reservoir. The compounds may also be tailored to optionally scavenge hydrogen sulphide produced by bacterial growth from the reservoir fluids.

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METHOD OF INHIBITING RESERVOIR SOURING BY BACTERIA 1 2 The present invention relates to a method of reducing 3 the souring of hydrocarbons due to bacterial production 4 of hydrogen sulphide gas. 5 6 During the production of hydrocarbon fluids, such as 7 oil or gas, it is common practice to flood the 8 9 reservoir holding the hydrocarbons with water to enhance production by maintaining pressure as oil 10 and/or gas is removed and to "sweep" hydrocarbons which 11 would normally remain within the reservoir to the 12 collection well. 13 14 The source water used in such flooding operations is 15 frequently a natural brine and in the case of offshore 16 oil and gas production is normally sea water. 17 18 Flooding of a reservoir with natural brines usually has 19 20 undesirable consequences since the brine may be responsible for the introduction and/or accelerated 21 22 growth of bacteria within a reservoir as follows: 23 24 Bacteria within the brine flood water are 1. introduced to the reservoir and find a hydrocarbon 25

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| L | source (for example, oil). | The oil may then be |
|---|------------------------------|---------------------|
| 2 | metabolised by the bacteria | allowing the rapid |
| 3 | growth of bacterial populat: | ions. |

4 5

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7 8 2. Bacteria which may be naturally present in the reservoir in dormant form will receive a nutrient supply due to the introduction of the flood water. Again, the increased nutrient supply will promote bacterial growth within the reservoir.

9 10

Generally, anaerobic conditions will exist in the 11 12 reservoir, and bacteria able to cope with low amounts 13 of oxygen or the absence of oxygen will preferentially 14 proliferate. Generally, the most dominant form of 15 bacteria is the so called sulphate reducing bacteria 16 (SRBs). SRBs metabolise sulphates from the injected flood water and produce hydrogen sulphide gas as a 17 waste product. 18

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The hydrogen sulphide generated will become partitioned in both the oil/gas and water phases within the reservoir. More importantly, however, the hydrogen sulphide would normally be co-produced in the fluids at the producing well.

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The process of hydrogen sulphide production by bacteria is known as reservoir souring. The consequence of this process is a gradual increase in hydrogen sulphide content within the produced fluids.

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Hydrogen sulphide is highly undesirable in the produced fluids since it is corrosive and toxic. Consequently, the presence of hydrogen sulphide as a contaminant adversely affects the sales quality of the exported oil and gas.

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It is thus normal practice within the industry to 1 attempt to prevent bacterial growth within the 2 reservoir by the addition of biocides to the injected 3 water in order to sterilise the system and therefore 4 prevent proliferation of bacteria. 5 6 Typical biocides include chlorine, aldehydes such as 7 gluteraldehde, thiazolines and quaternary amines. 8 9 However, the continual addition of biocide is 10 frequently too costly to practice and more usually a 11 sweep dosage at suitable intervals is used to sterilise 12 13 the system. 14 Despite biocide addition, it is almost impossible to 15 prevent the introduction of all bacteria to a reservoir 16 and a reduced rate of souring may still occur. 17 18 Additionally, due to the surface active nature of the 19 majority of biocides, the biocides are readily absorbed 20 to the surfaces of minerals such as clays found in 21 reservoir sands. It is therefore not normally possible 22 to achieve penetration of the biocide deep into a 23 reservoir. 24 25 Hence elimination of bacteria naturally present within 26 a reservoir or introduced by the flood water and moving 27 with oil/water flood front boundary is normally not 28 achievable using conventional biocides. 29 30 One way to obtain an end product having acceptably low 31 levels of hydrogen sulphide is the removal of that gas 32 by treating the extracted hydrocarbons. This will 33 normally require the installation of process equipment 34 to remove the hydrogen sulphide gas by, for example, 35 flaring, treating with amine scrubbers to absorb the 36

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gas or addition of chemicals to scavenge by reaction to produce a neutralised compound.

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However, all such methods of hydrogen sulphide removal considerably increase the cost of production and are generally considered to be undesirable.

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In summary, it can be seen that a number of options
exist to prevent build up of hydrogen sulphide in the
produced fluids:

11

12 a. inhibit the growth of bacteria within the13 reservoir by chemical addition to the flood water;

14

b. scavenge the evolved hydrogen sulphide within the
 reservoir to prevent production with recovered
 fluids.

18

However, as indicated above, it is apparent that such treatments are not practical using conventional biocides or scavengers from performance and/or cost constraints.

23

We have now found that the use of peroxy compounds
(which are not normally thought of as biocides) are
ideally suited to the treatment of bacterially
contaminated reservoirs. The peroxy compounds can both
destroy resident bacteria and also act to remove
hydrogen sulphide already generated.

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In one aspect, the present invention provides a method of inhibiting the growth of bacteria in a hydrocarbon reservoir, said method comprising introducing a peroxy compound to said reservoir.

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36 In another aspect, the present invention provides a

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method of combatting the generation of hydrogen 1 sulphide in a hydrocarbon reservoir by sulphur reducing 2 bacteria, said method comprising introducing a peroxy 3 compound to said reservoir. 4 5 The complementary actions of the peroxy compounds arise 6 from the oxidising potential of such compounds and the 7 generation of free radicals as the peroxy compound 8 9 decomposes. 10 The peroxy compounds for use in the present invention 11 are normally stable neutral compounds which therefore 12 show minimal absorption to the reservoir minerals and 13 can therefore penetrate deep into the reservoir carried 14 by the flood water. 15 16 The peroxy compounds of interest may be selected to 17 decompose at the particular temperatures encountered 18 within the reservoir (usually elevated temperatures 19 relative to the ambient). Particular activation 20 temperatures can be engineered by selection and/or 21 modifications of the peroxy compounds used, to allow 22 the selective placement of compounds within reservoirs 23 of widely differing temperatures. 24 25 Decomposition of the peroxy compounds leads to the 26 generation of free radicals which are highly aggressive 27 The free radicals cause damage towards living cells. 28 and ultimately destroy cells such as bacteria. 29 30 In addition, the generation of coincident oxidising 31 species occurs and these oxidising species will react 32 with hydrogen sulphide leading ultimately to formation 33 of harmless sulphates in the produced fluids. 34 35

36 The peroxy compounds therefore have the advantages of

| 1 | being selective to the destruction of bacteria without |
|----|--|
| 2 | being lost to the reservoir by absorption or |
| 3 | decomposition prior to deep penetration. The peroxy |
| 4 | compounds also remove encountered hydrogen sulphide to |
| 5 | prevent continued production within the recovered |
| 6 | fluids. |
| 7 | |
| 8 | The type of peroxy compounds useful for this invention |
| 9 | contain the chemical grouping -0-0- and may be |
| 10 | generally described by the formula: |
| 11 | |
| 12 | R-O-O-R' |
| 13 | |
| 14 | where R and R' may be hydrogen, alkyl or aromatic |
| 15 | groups (including substituted cyclic and branched |
| 16 | groups), oxygenated hydrocarbon chains such as |
| 17 | carbonate or ketone, or other groups which will result |
| 18 | in stabilisation of the peroxide linkage. |
| 19 | |
| 20 | Modifications to the structure of R and R' alter the |
| 21 | solubility characteristics and thermal stabilities of |
| 22 | the peroxides which allows for selective placement and |
| 23 | activation in reservoirs of differing temperatures and |
| 24 | can also allow penetration of oil layers by selection |
| 25 | of oleophilic peroxides should this be desirable. |
| 26 | |
| 27 | The reaction occurring by thermal decomposition of the |
| 28 | peroxy compounds leads to the generation of free |
| 29 | radicals, thus: |
| 30 | Δн |
| 31 | $R'-O-O-R \rightarrow R'-O \cdot + R-O \cdot$ |
| 32 | |
| 33 | The generated free radicals may undergo various |
| 34 | reactions leading to complex organic products but all |
| 35 | such reactions lead to cellular attack and therefore |
| 36 | have biocidal properties. |

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In addition, the reaction: 1 2 $R-O \bullet + H_2S \rightarrow R-OH + HS \bullet$ 3 4 can take place to give a reactive thiol free radical 5 (HS.) which will bind to organic materials such as 6 crude oil or bacterial cells to eliminate free H_2S from 7 the system. 8 9 A further decomposition reaction can lead to the 10 formation of free oxygen which again limits bacterial 11 activity by altering the oxygen free conditions 12 necessary for SRB growth. 13 14 The described peroxy compounds therefore provide 15 advantageous multifunctional behaviour which can 16 prevent reservoir souring. 17 18 Viewed from a further aspect, the present invention 19 provides the use of a peroxy compound to inhibit 20 bacterial growth within a hydrocarbon reservoir or 21 to reduce the hydrogen sulphide content of a 22 hydrocarbon fluid. 23 24 In addition, the present invention also provides a 25 hydrocarbon fluid (eg oil or gas) containing a peroxy 26 compound or a reactant thereof as an additive. 27 28 The range of useful peroxy compounds to be applied to a 29 reservoir may be characterised by a self accelerating 30 decomposition temperature (SADT) where a critical value 31 will cause the compound to decompose at an accelerating 32 rate yielding the free radicals necessary for biocidal 33 34 activity. 35

36 However, the value may be adjusted by the inclusion of

1 accelerators or inhibitors which allow a peroxy 2 compound to be utilised at temperatures above or below 3 this figure. 4 5 Examples of suitable peroxy compounds which may be 6 added to a reservoir flood water for the purpose of the 7 invention include, but are not limited to: 8 9 Methyl ethyl ketone peroxide, 10 Cyclohexanone peroxide, 11 Acetyl acetone peroxide, 12 Diacetone alcohol peroxide, 13 Dibenzoyl peroxide, 14 Ditertiary butyl peroxide, 15 Tertiary butyl peroxide, 16 Hydrogen peroxide, 17 Tertiary butyl peroxy benzoate, 18 Inorganic and organic peroxycarbonates, or 19 20 functional equivalents or derivatives of these 21 compounds. A mixture of such compounds may be used. 22 23 Other compounds which are known to generate oxidative 24 free radicals by thermal or catalytic decomposition 25 will show similar biocidal activity. 26 27 Generally, the peroxy compounds will be added to the 28 flood water prior to injection into the reservoir.

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1 CLAIMS

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1. A method of inhibiting the growth of bacteria in a hydrocarbon reservoir, said method comprising introducing a peroxy compound to said reservoir.

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7 2. A method of combatting the generation of hydrogen 8 sulphide in a hydrocarbon reservoir by sulphur 9 reducing bacteria, said method comprising 10 introducing a peroxy compound to said reservoir.

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12 3. A method as claimed in either one of Claims 1 and
13 2, wherein the peroxy compound may consist of a
14 water soluble or water dispersable inorganic or
15 organic based peroxy compound.

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17 4. A method as claimed in any one of Claims 1 to 3,
18 wherein the peroxy compound may be a peroxide,
19 hydroperoxide, peroxycarbonate, peroxysulphate or
20 other peroxide based complex.

21

22 5. A method as claimed in any one of Claims 1 to 4, wherein the peroxy compound may be selected from 23 24 one or more of methyl ethyl ketone peroxide, cyclohexanone peroxide, acetyl acetone peroxide, 25 26 diacetone alcohol peroxide, dibenzoyl peroxide, ditertiary butyl peroxide, tertiary butyl 27 peroxide, hydrogen peroxide, tertiary butyl peroxy 28 29 benzoate, inorganic and organic peroxycarbonates, 30 or functional equivalents or derivatives of these 31 compounds.

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33 6. A method as claimed in any one of Claims 1 to 5, 34 wherein said peroxy compound is added to the 35 reservoir flood water.

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| 1 | 7. | A method as claimed in any one of Claims 1 to 6, |
|----|----|--|
| 2 | | wherein the peroxy compound is selected to |
| 3 | | decompose at a temperature of the reservoir. |
| 4 | | |
| 5 | 8. | A method as claimed in any one of Claims 1 to 7, |
| 6 | | wherein a co-additive may be optionally added to |
| 7 | | the peroxy compound to either inhibit or |
| 8 | | accelerate decomposition of the peroxy compound as |
| 9 | | a function of temperature. |
| .0 | | |

10 11

9. Use of a peroxy compound to inhibit bacterial growth within a hydrocarbon reservoir.

12 13

14 10. Use of a peroxy compound to reduce the hydrogen 15 sulphide content of a hydrocarbon fluid.

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17 11. Hydrocarbon fluid containing a peroxy compound or a reactant thereof as an additive.

INTERNATIONAL SEARCH REPORT

Interna .ial application No. PCT/GB 97/01537

| A. CLASSIFICATION OF SUBJECT MATTER IPC6: C106 27/12, A01N 37/00 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC6: C106, A01N Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Bectronic data base controlled during the international search (name of data base and, where practicable, search terms used) WPI, APIPAT C. DOCUMENTS CONSIDERED TO BE RELEVANT Category* Citation of document, with indication, where appropriate, of the relevant passages X. AU 53138/79 A1 (BIOLEX CORP.), 28 May 1981 (28.05.81), page 6, line 35 - page 8, line 1 - line 16, claims 1-7 A EP 0082513 A1 (PEROXID-CHEMIE GMBH), 29 June 1983 1-11 Purchase documents are listed in the continuation of Box C. Special categories of citoud documents **Occounts with may have documents are listed in the continuation of Box C. **Occounts defining the greend stee of the set which is not considered the set and not in contile with the application documents of the continuation of Box C. **Occounts of publication and an order citations or other than the principle of citoud documents. A continuation of the international filing date or principle to an out disclosure, use, exhibition or other citations | | | 707/02 37,0 | |
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INTERNATIONAL SEARCH REPORT

Information on patent family members

06/08/97

International application No.
PCT/GB 97/01537

| Publication date | Patent family member(s) | Publication date |
|------------------|--|---|
| 28/05/81 | NONE | |
| 29/06/83 | CA 1205616 A DE 3151133 A US 4548708 A | 10/06/86 30/06/83 22/10/85 |
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